Amendments to the Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims:

- 1. (Original) Isolated protein comprising the same or substantially the same amino acid sequence selected from the group consisting of SEQ ID NOs: 1 to 12, or a splice variant or a salt thereof.
- 2. (Original) Protein according to claim 1, which comprises at least one fragment of said amino acid sequences.
- 3. (Currently Amended) Nucleic acid, which comprises at least one nucleic acid encoding a protein according to claims 1-and/or-2.
- 4. (Original) Nucleic acid according to claim 3, which consists of DNA or RNA.
- 5. (Original) Nucleic acid according to claim 4 comprising a nucleic acid selected from the group consisting of SEQ ID NOs: 13 to 24.
- 6. (Currently Amended) Nucleic acid according to one of claims 3 to 5claim 3 further comprising at least one promoter, enhancer, intron and/or polyAsequence.
- 7. (Currently Amended) Nucleic acid, which is complementary to the nucleic acid according to one of claims 3 to 6claim 3.
- 8. (Currently Amended) Vector comprising a protein according to claims 1 or 2 and/or a nucleic acid according to one of claims 3 to 7claim 1.

- 9. (Original) Vector according to claim 8 selected from the group of vectors consisting of plasmids, phagemids, phages, cosmids, artificial mammalian chromosomes, knock-out or knock-in constructs, viruses, in particular adenovirus, vaccinia virus, baculovirus, retrovirus, adeno-associated-virus, rhinovirus, HIV, adeno-associated virus (AAV), herpes simplex virus (HSV-1), lentivirus, filovirus and engineered versions thereof, naked DNA, virosomes, liposomes, nucleic acid coated particles, in particular gold spheres.
- 10. (Currently Amended) Isolated cell comprising a protein according to claim 1claims 1 or 2, a nucleic acid according to claims 3 to 7 and/or a vector according to claims 8 or 9.
- 11. (Original) Cell according to claim 10, which is a stem cell, a neuronal precursor cell or a neuronal cell, in particular an axon.
- 12. (Currently Amended) Transgenic non-human animal generated from a cell according to claims 10 or 1110.
- 13. (Currently Amended) Antibody directed against a protein according to claims 1 or 2claim 1.
- 14. (Currently Amended) Method of producing a protein comprising the same or substantially the same amino acid sequence selected from the group consisting of SEQ ID NOs: 1 to 12, or a splice variant or a salt thereofaccording to claims 1 or 2 and/or a nucleic acid according to claims 3 to 7 comprising the steps of:
 - a) cultivating a cell according to claim 10-or-11, and
 - b) isolating said protein and/or said nucleic acid.

- 15. (Currently Amended) Method of isolating compounds interacting with a protein according to claims 1 or 2 claim 1 comprising the steps of:
 - a) contacting said protein with at least on potentially interacting compound,
 - b) measuring binding of said compound to said protein.
- 16. (Original) Method according to claim 16, further comprising the steps of:
 - a) selecting a binding compound,
 - b) modifying the binding compound, to generate a variety of modified binding compounds,
 - c) contacting said protein with each of the modified binding compounds,
 - d) measuring binding of said modified compounds to said protein, and
 - e) if needed repeating steps a) to d) for one or more times.
- 17. (Original) Method of isolating functional interactors comprising the steps of:
 - a) contacting a neuronal cell that comprises a wt nucleic acid coding for a protein selected from the group consisting of SEQ ID NOs: 1 to 12, a splice variant thereof, or a fragment thereof with a potential functional interactor,
 - b) contacting the cell with a bioactive lipid phosphate, and
 - c) measuring neurite movement.
- 18. (Original) Method according to claim 17 further comprising the steps of:
 - a) contacting a neuronal cell that comprises a mutant nucleic acid coding for a
 mutant of the protein selected from the group consisting of SEQ ID NOs: 1 to
 12, or a splice variant thereof or that contains a knock-out of the wt nucleic
 acid coding for one of said proteins with a potential functional interactor,
 - b) contacting said cell with a bioactive lipid phosphate, and

- c) measuring neurite movement.
- 19. (Currently Amended) Method according to claims 17 or 18 claim 17 further comprising the steps of:
 - a) selecting a functional interactor,
 - b) modifying the functional interactor, to generate a variety of modified functional interactors,
 - c) contacting a said neuronal cell as used in claim 17 a) and if needed a cell as used in claim 18 a) with each of the modified functional interactors,
 - d) contacting said cell or cells with a bioactive lipid phosphate,
 - e) measuring neurite movement, and
 - f) if needed repeating steps a) to d) for one or more times.
- 20. (Currently Amended) Method according to one of claims 15 to 19claim 15, further comprising the step of admixing the interacting compound or the functional interactor with suitable auxiliary substances and/or additives.
- 21. (Currently Amended) Pharmaceutical composition for the treatment of neuronal injuries or diseases comprising a protein according to claims 1 or 2, a nucleic acid according to claims 3 to 7, a vector according to claims 8 or 9, a cell according to claims 10 to 11, an antibody according to claims 13, a binding compound isolated by the method of claims 15 or 16 and/or a functional interactor isolated by the method of claims 17 to 19 claim 1 and if needed suitable auxiliary substances and/or additives.
- 22. (Original) Use of a pharmaceutical composition of claim 21 for the production of a medicament for the treatment of neuronal diseases or injuries, including spinal card lesions, head traumata, Alzheimer disease and stroke.

- 23. (Currently Amended) Use of a protein according to claim 1 or 2 or a nucleic acid according to claims 3 to 7 as a diagnostic marker for the diagnosis of a disease or disease state.
- 24. (Original) Use according to claim 23, where the disease is a neuronal disease, a tumor disease or infertility.
- 25. (New) Vector comprising a nucleic acid according to claim 3.
- 26. (New) Isolated cell comprising a nucleic acid according to claim 3.
- 27. (New) Isolated cell comprising a vector according to claim 8.
- 28. (New) Use of a nucleic acid of claim 3 as a diagnostic marker for the diagnosis of a disease or disease state.
- 29. (New) Use according to claim 29, where the disease is a neuronal disease, a tumor disease or infertility.

Amendments to the Sequence Listings:

Please insert the attached 59 page Sequence Listing into the application.